Supporting Information

Facile fabrication of composite electrospun nano-fibrous matrices of poly(εcaprolactone)-silica based Pickering emulsion

Archana Samanta^a, Sonam Takkar^b, Ritu Kulshreshtha^b, Bhanu Nandan^a, Rajiv K. Srivastava*

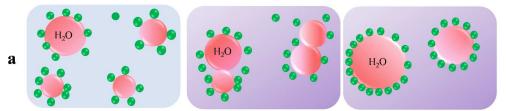
^a Department of Textile Technology, Indian Institute of Technology Delhi, Hauz Khas, New Delhi 110016, India

^b Department of Biochemical Engineering and Biotechnology, Indian Institute of Technology Delhi, Hauz Khas, New Delhi 110016, India

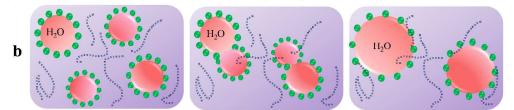
*Corresponding Author, Email: rajiv@textile.iitd.ac.in

Table of contents

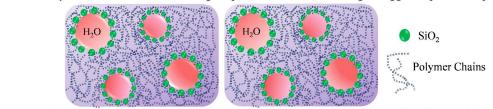
- 1. Scheme S1. Effect of m-silica and polymer concentration on emulsion
- 2. Figure S1(a): Visual observation of colloids (i) m-silica in toluene, (ii) m-silica in water and (iii) m-silica in 1:1 volume mixture of toluene and water
- 3. Figure S1 (b): Pink: Water + Rhodamine B and Transparent: PCL in toluene
- 4. Figure S2: Formation of stable water-in-oil emulsions of PCL using m-silica as the Pickering agent
- 5. Figure S3: Cryo-fractured cross-sections of PCL/m-silica emulsion films stabilized with 3wt.% or 5wt.% m-silica
- 6. Figure S4: Optical microscopic images of m-silica stabilized emulsions with maximum volume fraction of aqueous phase reflecting the reduction in droplet size with increase in polymer concentration
- 7. Figure S5: Variation of R^* (Ratio of surface area of n_0 number of m-silica particles per unit water-oil interfacial area) with respect to PCL and m-silica concentration
- Figure S6: SEM images of PCL/m-silica electrospun composite matrices made using 3 wt.% m-silica stabilized emulsions
- Figure S7: SEM images of PCL/m-silica electrospun composite matrices made using 3 or 5 wt.% m-silica stabilized emulsions with maximum volume fraction of aqueous phase
- 10. Figure S8: Effect of inclusion of m-silica on melting behaviour of PCL matrix
- 11. Figure S9: SEM images reflecting stability of PCL and PCL silica composite electrospun matrices to aqueous environment
- 12. Figure S10: Microscopic images showing fibroblast (L929) cell growth in different PCL/m-silica composite matrices in comparison to a matrix made using neat PCL



a) Reduced silica concentration at interface resulting in droplet coalescence and emulsion destabilization



b) Reduced Polymer Concentration favouring droplet coalescence, resulting in bigger dispersed droplets



С

c) Increased Polymer Concentration prevents droplet coalescence, favouring smaller dispersed droplets

Scheme S1: Effect of m-silica and polymer concentration on emulsion stability (a) low concentration of Pickering stabilizer leading to emulsion destabilization (b) reduced polymer concentration leading to droplet coalescence (c) higher polymer concentration preventing droplet coalescence

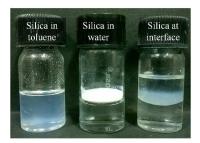
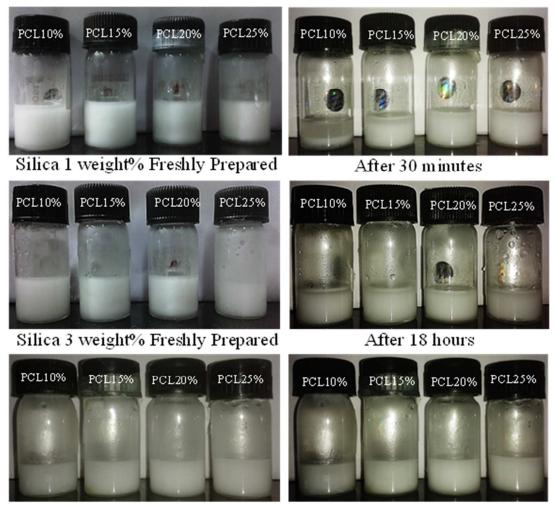


Figure S1 (a): Visual observation of colloids (i) m-silica in toluene, (ii) m-silica in water and (iii) m-silica in 1:1 volume mixture of toluene and water



Figure S1 (b): Pink: Water + Rhodamine B and Transparent: PCL in toluene



Silica 5 weight% Freshly Prepared

After 50 hours

Figure S2: Formation of stable water-in-oil emulsions of PCL using m-silica as the Pickering agent

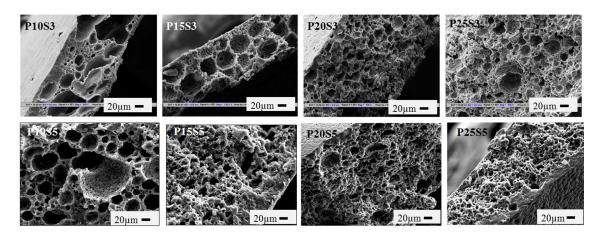


Figure S3: Cryo-fractured cross-sections of PCL/m-silica emulsion films stabilized with 3wt.% or 5wt.% m-silica

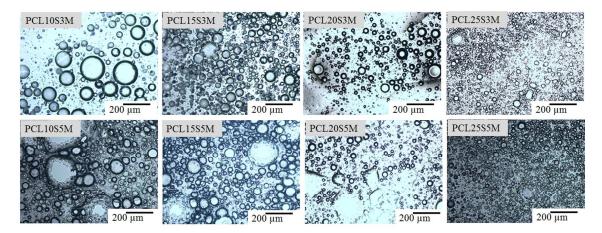


Figure S4: Optical microscopic images of m-silica stabilized emulsions with maximum volume fraction of aqueous phase reflecting the reduction in droplet size with increase in polymer concentration

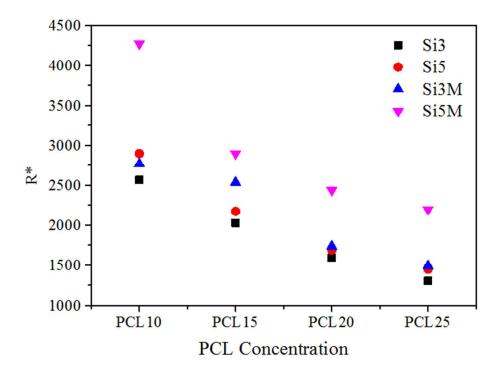


Figure S5: Variation of R* (Ratio of surface area of n₀ number of m-silica particles per unit water-oil interfacial area) with respect to PCL and m-silica concentration

Si3 and Si5 represents samples stabilized with 3 and 5 weight percent silica with respect to oil phase respectively, with $\phi_d=0.26$ against corresponding PCL concentration as denoted on the x-axis.

Si3M and Si5M represents samples stabilized with 3 and 5 weight percent silica with respect to oil phase respectively, with maximum ϕ_d against corresponding PCL concentration as denoted on the x-axis

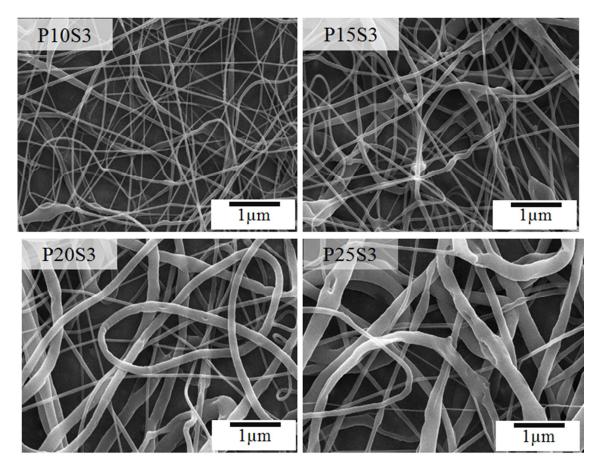


Figure S6: SEM images of PCL/m-silica electrospun composite matrices made using 3 wt.% m-silica stabilized emulsions

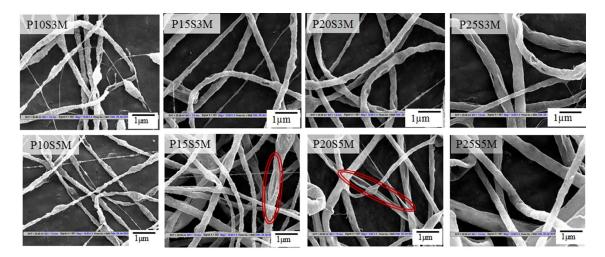


Figure S7: SEM images of PCL/m-silica electrospun composite matrices made using 3 or 5 wt.% m-silica stabilized emulsions with maximum volume fraction of aqueous phase

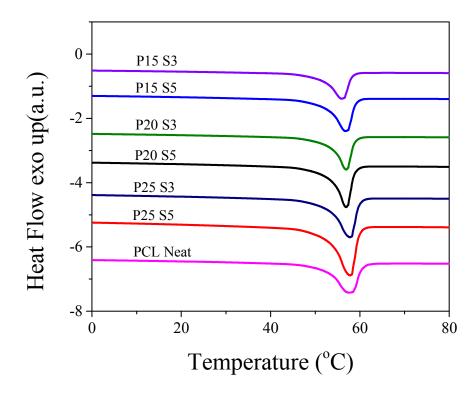
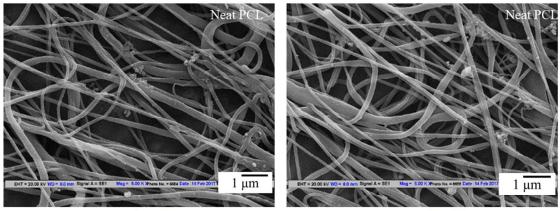


Figure S8: Effect of inclusion of m-silica on melting behaviour of PCL matrix



Day 1

Day 21

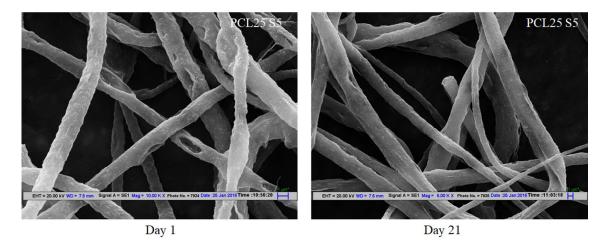


Figure S9: SEM images reflecting stability of PCL and PCL silica composite electrospun matrices to aqueous environment

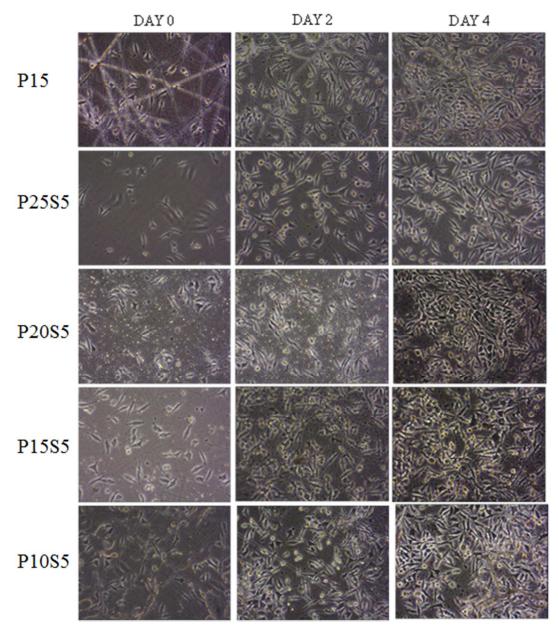


Figure S10: Microscopic images showing fibroblast (L929) cell growth in different PCL/msilica composite matrices in comparison to a matrix made using neat PCL